

Article

EPI-001, a compound active against castration-resistant prostate cancer, targets transactivation unit 5 of the androgen receptor

Eva De Mol, R. Bryn Fenwick, Christopher T.W. Phang, Victor Buzon, Elzbieta Szulc, Alex De La Fuente, Albert Escobedo, Jesús García, Carlos W. Bertoncini, Eva Estebanez-Perpina, Iain J. McEwan, Antoni Riera, and Xavier Salvatella

ACS Chem. Biol., **Just Accepted Manuscript** • DOI: 10.1021/acscchembio.6b00182 • Publication Date (Web): 29 Jun 2016

Downloaded from <http://pubs.acs.org> on June 30, 2016

Just Accepted

"Just Accepted" manuscripts have been peer-reviewed and accepted for publication. They are posted online prior to technical editing, formatting for publication and author proofing. The American Chemical Society provides "Just Accepted" as a free service to the research community to expedite the dissemination of scientific material as soon as possible after acceptance. "Just Accepted" manuscripts appear in full in PDF format accompanied by an HTML abstract. "Just Accepted" manuscripts have been fully peer reviewed, but should not be considered the official version of record. They are accessible to all readers and citable by the Digital Object Identifier (DOI®). "Just Accepted" is an optional service offered to authors. Therefore, the "Just Accepted" Web site may not include all articles that will be published in the journal. After a manuscript is technically edited and formatted, it will be removed from the "Just Accepted" Web site and published as an ASAP article. Note that technical editing may introduce minor changes to the manuscript text and/or graphics which could affect content, and all legal disclaimers and ethical guidelines that apply to the journal pertain. ACS cannot be held responsible for errors or consequences arising from the use of information contained in these "Just Accepted" manuscripts.



ACS Publications

ACS Chemical Biology is published by the American Chemical Society, 1155 Sixteenth Street N.W., Washington, DC 20036

Published by American Chemical Society. Copyright © American Chemical Society. However, no copyright claim is made to original U.S. Government works, or works produced by employees of any Commonwealth realm Crown government in the course of their duties.

**EPI-001, a compound active against castration-resistant prostate cancer, targets
transactivation unit 5 of the androgen receptor**

Eva De Mol,¹ R. Bryn Fenwick,¹ Christopher T. W. Phang,¹ Victor Buzón,² Elzbieta Szulc,¹ Alex de la Fuente,¹ Albert Escobedo,¹ Jesús García,¹ Carlos W. Bertoncini,¹ Eva Estébanez-Perpiñá,² Iain J. McEwan,³ Antoni Riera^{1,4} and Xavier Salvatella^{1,5,*}

¹Institute for Research in Biomedicine (IRB Barcelona), The Barcelona Institute of Science and Technology, Baldiri Reixac 10, 08028 Barcelona, Spain.

²Departament de Bioquímica i Biologia Molecular, Universitat de Barcelona and Institute of Biomedicine of the University of Barcelona (IBUB), Baldiri Reixac 10, 08028 Barcelona, Spain.

³Institute of Medical Sciences, School of Medicine, Medical Sciences and Nutrition, University of Aberdeen, IMS Building, Foresterhill, Aberdeen AB25 2ZD, Scotland, United Kingdom.

⁴Departament de Química Orgànica, Universitat de Barcelona, Martí i Franqués 1-11, 08028 Barcelona, Spain.

⁵ICREA, Passeig Lluís Companys 23, 08010 Barcelona, Spain.

*to whom correspondence should be addressed: +34 934020459,
xavier.salvatella@irbbarcelona.org.

1
2
3
4
5 **Castration-resistant prostate cancer is the lethal condition suffered by prostate**
6 **cancer patients that become refractory to androgen deprivation therapy. EPI-001**
7 **is a recently identified compound active against this condition that modulates the**
8 **activity of the androgen receptor, a nuclear receptor that is essential for disease**
9 **progression. The mechanism by which this compound exerts its inhibitory activity**
10 **is however not yet fully understood. Here we show, by using high resolution**
11 **solution nuclear magnetic resonance spectroscopy, that EPI-001 selectively**
12 **interacts with a partially folded region of the transactivation domain of the**
13 **androgen receptor, known as transactivation unit 5, that is key for the ability of**
14 **prostate cells to proliferate in the absence of androgens, a distinctive feature of**
15 **castration-resistant prostate cancer. Our results can contribute to the**
16 **development of more potent and less toxic novel androgen receptor antagonists**
17 **for treating this disease.**
18
19
20
21
22
23
24
25
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60

INTRODUCTION

Prostate cancer (PC) is the second most common cancer in men and can be cured by surgery or radiotherapy in ca 70% of cases. The first line of pharmacological treatment for the remaining cases targets the androgen receptor (AR) because prostate cells depend on its activation by androgens for their growth and proliferation¹. Activation can be prevented by combining androgen deprivation therapy, which inhibits the secretion of androgens by the testes, with the administration of antagonists that competitively bind to the binding site of androgens in the ligand-binding domain (LBD) of AR².

Two to three years into this treatment PC patients inevitably develop castration-resistant prostate cancer (CRPC) as prostate cancer cells acquire the ability to activate AR at low levels of circulating androgens and in the presence of antagonists³. The mechanisms of aberrant activation are not well understood but appear to include the amplification of the AR gene and AR overexpression, the expression of constitutively active AR splice variants lacking the LBD, cell signaling cross-talk and mutations in both AR and transcriptional co-regulators⁴.

AR is a large multi-domain protein composed of globular ligand- and DNA-binding domains (LBD and DBD) and an N-terminal transactivation domain (NTD) that is intrinsically disordered (ID)^{5,6} (Fig. 1a). The function of the NTD (residues 1 to 558) is to recruit the basal transcription machinery by binding to general transcription factors either directly or assisted by transcriptional co-activators¹. These protein-protein interactions are thought to cause the folding of binding motifs in a region of the NTD called activation function 1 (AF-1) that has not yet been characterized at high resolution (Fig. 1a,b)⁶. Inhibiting these interactions is considered a potential therapeutic approach for both PC and CRPC⁷, but the NTD has not been considered a suitable target for drug discovery due to its apparent lack of persistent secondary and tertiary structure.

The development of drugs that interact with ID regions has however recently been met with some success^{8,9}, and has shown that targeting them with small molecules may be a viable therapeutic approach^{10,11}. A particularly important development in this area was the recent discovery of EPI-001, an experimental drug for the treatment of CRPC identified by phenotypic screening that is efficacious both in cell lines and in an animal

model of this disease¹². *In vivo* EPI-001 binds irreversibly to the AR NTD and weakens its interaction with general transcription factors and transcriptional co-activators¹³.

The discovery of EPI-001 represents an important milestone and a derivative of this compound is currently in clinical trials for CRPC (NCT02606123). The lack of a detailed understanding of the structural features of the domain and of the mechanism of action of this class of compounds represents, however, a hurdle for the rational design of optimized inhibitors. Here we reveal, by using solution nuclear magnetic resonance (NMR) spectroscopy, the structural properties of the ID regions of the NTD predicted to have a high propensity to fold and show that EPI-001 targets a region of sequence, known as transcription activation unit 5 (Tau-5), that is key for the ability of prostate cells to proliferate in the absence of androgens.

METHODS

Protein expression and purification The DNA sequences coding for human WT AR residues 265 to 340 (AF-1*₂₆₅₋₃₄₀), 330 to 448 (Tau-5*) and 142 to 448 (AF-1*) were cloned into Gateway pDEST17 vectors (Invitrogen) with an N-terminal His₆-tag and a TEV cleavage site. Transformed *E. coli* Rosetta cells were grown at 37°C in LB medium for the production of non-isotopically labeled protein. For single (¹⁵N) or double (¹⁵N, ¹³C) isotopic labeling, cells were grown in minimal MOPS medium¹⁴ containing ¹⁵NH₄Cl or ¹⁵NH₄Cl and ¹³C-glucose, respectively. The AF-1* fusion protein accumulated in inclusion bodies which were solubilized in lysis buffer containing 8 M urea and fragmented by a pass through a cell disruptor at 25 kpsi. The fusion protein was purified by Ni²⁺ affinity chromatography in urea, which was removed by two dialysis steps, after which the His₆-tag was cleaved by the TEV protease. The cleaved AF-1* was further purified by reverse Ni²⁺ affinity and size exclusion chromatography in 20 mM sodium phosphate buffer with 1 mM tris(2-carboxyethyl)phosphine hydrochloride (TCEP) at pH 7.4.

Peptides The synthesis of peptides R1, R2 and R3 (See SI) was performed by solid phase peptide synthesis by GenScript (peptide R1) or by ICTS NANBIOSIS, more specifically by the peptide synthesis unit of the CIBER in bioengineering, biomaterials & nanomedicine (CIBER-BBN) at the Barcelona science park (peptides R2 and R3). The

lyophilized peptides were dissolved in deionized water and the pH of the resulting solution was adjusted by addition of a concentrated NaOH solution. The concentration of these solutions was determined by amino acid analysis. The absence of intermolecular disulfide bonds in the R2 peptide, which contains one Cys residue, was confirmed by mass spectrometry (MS).

Chemical synthesis of EPI-001 and stereoisomers EPI-001 contains two stereogenic centers and can therefore be found as two pairs of enantiomers. We synthesized the four isomers with high diastereo- and enantioselectivity (Chiral HPLC). The synthesis followed the sequence detailed in Fig. S1 of the SI. Bisphenol A was treated with enantiomerically pure glycidol. The resulting diol was protected as dimethyl acetal and the free phenol was allowed to react with another isomer of glycidol. The corresponding diol was transformed into the epoxide and opened with CeCl_3 . The treatment gave the final product by concomitant deprotection of the acetal. Full experimental details as well as the complete characterization of all isomers can be found in the SI.

NMR The assignment of AF-1* was obtained by using a *divide and conquer* approach. The resonances of fragments AF-1*₂₆₅₋₃₄₀, Tau-5* (330 to 448) and AF-1* (142-448) were obtained by analyzing conventional three-dimensional triple resonance experiments acquired with standard Bruker pulse sequences on Bruker 600 MHz and 800 MHz spectrometers at 278 K in 20 mM sodium phosphate buffer with 1 mM TCEP at pH 7.4. The resonances of AF-1*₂₆₅₋₃₄₀ and Tau-5* were equivalent to those of AF-1*, except for residues near the termini (Fig. S2), and allowed transferring the assignments from the former to the latter; the residues that were unique to AF-1* were assigned directly by analysis of the relevant spectra. To measure the perturbations caused by EPI-001, EPI-002, EPI-003, EPI-004 and EPI-005 (details available as SI) on the resonances of AF-1* appropriate volumes of a 50 mM stock solution of these compounds in 100% dioxane- d_8 were added to aliquots containing 25 μM AF-1*, 20 mM sodium phosphate, 1 mM TCEP, 10% D_2O , 30 μM DSS- d_6 at pH 7.4.

RESULTS AND DISCUSSION

As previously reported^{5,6,15} the sequence of the NTD has features typically encountered in ID regions¹⁶. It has a high content of Gly, Pro, polar and charged residues and, as

shown in figure 1b, possesses regions of low sequence complexity such as polyGln (residues 58-78, 84-89, 193-197), polyPro (372-379), polyAla (398-402) and polyGly (449-472) tracts. Such disordered tracts are commonly flanked by motifs of relatively low disorder propensity, with some helical propensity and rich in hydrophobic side chains, that are often involved in interactions with binding partners¹⁷. An analysis of the sequence of the NTD with predictors of disorder and helical propensity indicates that several such motifs are indeed present in the domain and it has been proposed that they may correspond to the regions of sequence recognized by general transcription factors and transcriptional co-regulators¹⁵ (Fig. 1c,d,e).

A number of studies have aimed at identifying the regions of sequence of AF-1 that are essential for transcriptional activity^{18,19}. Two large regions of sequence, known as transcription activation units 1 and 5 (Tau-1, 102-371 and Tau-5, 361-537) have emerged as especially important (Fig. 1b). Tau-1 is crucial for the transcriptional activity of AR when the receptor is activated by androgens¹⁸. Although Tau-5 is less well characterised, it has been shown to be more important than Tau-1 when activation occurs via androgen independent mechanisms in androgen depletion independent cell lines derived from CRPC patients and mouse xenograft models of PC^{20,21}. Efforts have also been directed at identifying the critical stretches of residues within these regions and suggest that for Tau-1 they correspond to residues 174 to 204 (core Tau-1)¹⁹, and for Tau-5 to residues 433 to 437 (WHTLF motif)²⁰ (Fig. 1c).

AF-1* is partially folded To investigate the structural properties of the NTD and its interaction with EPI-001 we cloned, expressed, purified and studied by NMR at 278K a 308-residue construct (AF-1*, residues 142-448) containing the part of AF-1 predicted to have a low disorder propensity (Fig. 1b and 1c), that is flanked by polyGln and polyGly tracts (Fig. 1b). In agreement with previous reports based on the use of other biophysical methods⁶ the ¹H,¹⁵N-HSQC spectrum of AF-1* had the features expected in an ID region such as low H^N chemical shift dispersion (Fig. 2a). In addition, and contrary to what is the case for NTD constructs containing residues 1 to 141, which include the polyQ tract (Fig. 1b), this region of the NTD was sufficiently stable to allow a characterization of its structural properties by solution NMR.

In spite of its ID nature, the AF-1 region of the NTD has been shown to have helical propensity by circular dichroism (CD) in buffer and in the presence of cosolvents that stabilize intramolecular hydrogen bonds, as well as in the presence of the natural osmolyte trimethylamine oxide (TMAO)^{5,22}. To identify the regions of sequence that adopt this secondary structure we assigned the resonances of AF-1* by a *divide and conquer* approach (see Methods), compared the C α and C β chemical shifts, which are reliable reporters of secondary structure, to those predicted for disordered AF-1* (Fig. 2b)²³ and used the SSP algorithm of Forman-Kay and co-workers²⁴ to quantify the secondary structure propensity for the various residues of this construct from analysis of the backbone (¹³C α , ¹³C β , ¹³CO, ¹⁵N, H^N) chemical shifts.

The results that we obtained indicated the presence of two regions of high helical propensity (defined as $\Delta\delta\text{C}\alpha\text{-}\Delta\delta\text{C}\beta > 1$ ppm and SSP ≈ 0.5 , corresponding to a helical propensity of 50%) which correspond to residues 185-200, in Tau-1, and 390-410, in Tau-5. Other regions of intermediate helical propensity (defined as $\Delta\delta\text{C}\alpha\text{-}\Delta\delta\text{C}\beta \approx 0.5$ ppm and SSP ≈ 0.2) could be also identified, such as the region 230-240, in Tau-1, and 355-365 in Tau-5 (Fig. 2b, c). In addition to identifying regions of helical secondary structure the analysis of the chemical shifts also suggests that residues 144-154 and 270-290 of AF-1* adopt an extended conformation ($\Delta\delta\text{C}\alpha\text{-}\Delta\delta\text{C}\beta \approx -0.5$ ppm and SSP ≈ -0.3) (Fig. 2b, c).

To further characterize the structural properties of AF-1* we characterized the dynamics of AF-1* by measuring the transverse relaxation rates (R_2) of the backbone ¹⁵N nuclei. Such relaxation rates are good reporters of nascent secondary structure and transient tertiary contacts in chemically denatured and ID proteins²⁵. The results that we obtained, presented in figure 2d, indicate that the regions with nascent secondary structure revealed by the analysis of the ¹³C chemical shifts (Fig. 2b) also display relatively high R_2 values. These are especially high, reaching values of *ca* 15 s⁻¹, for three regions of sequence found in Tau-5 predicted to have low disorder propensity²⁶ and presenting, in two cases, high helical propensity.

Some of the regions of sequence that we identified as partially folded in AF-1* correspond in fact to the epitopes of binding partners of AR or are known to be important for the function of this receptor. In Tau-1, for example, the region that is most structured corresponds to *core Tau-1* (Fig. 2b). This region encompasses the motif

¹⁸³LSEASTMQLL¹⁹², which is the binding epitope of TAB2, a component of the NCoR co-repressor complex²⁷. Partially folded residues 144 to 154, also in Tau-1, are part of the binding epitope of the amino-terminal bromodomains of BRD4²⁸. The region containing residues 230 to 240, of intermediate helicity, overlaps with the binding site of CHIP, a protein known to recruit the chaperone machinery by interaction with Hsc70, Hsp70 and Hsp90, which mediates the degradation of AR by the proteasome²⁹ (Fig. 1e).

In Tau-5 the regions that we found to be partially folded are separated by motifs of low sequence complexity such as the Pro-rich region ³⁷¹GPPPPPPPPHP³⁸¹ and the Gly-rich region ⁴¹⁴GAGAAGPGSGSPS⁴²⁶. The former has been proposed to be the binding site of the SH3 domain of Src³⁰ and the latter harbors a phosphosite, S424, which is thought to be important for transcriptional activity³¹. Although we have not assigned the backbone resonances corresponding to the Pro residues in this Pro-rich motif, an analysis of the secondary chemical shifts of residues in its flanking regions (367-370, 385-388, Fig. 2b) suggests that it may adopt an extended conformation, likely of the polyproline II type, given the high Pro content; the polyproline II conformation is rather unusual but has been documented for several other IDPs.³² The chemical shifts and transverse ¹⁵N relaxation rates of the Gly-rich region (414-426, Fig. 2b,c,d) indicate, by contrast, that it is disordered, as expected due to its high Gly content.

No binding partners have yet been identified for regions 355-365 and 390-410, that are partially helical, but the ⁴³³WHTLF⁴³⁷ motif in region 433-447, corresponding to the core of Tau-5, has been proposed to bind both to activation function 2 (AF-2) in the LBD of androgen-bound AR^{33,34} and to histone acetyltransferase p300³⁵; in addition it has been shown to be indispensable for transcriptional activity in androgen depletion independent cell lines²⁰.

We noted that the helical propensity of the partially folded regions identified in both Tau-1 and Tau-5 (SSP \approx 0.5, corresponding to a population of helix of ca 50%) is substantially higher than that predicted by algorithms that solely consider local interactions such as Agadir³⁶ (Figs. 1e and 2b, c), indicating that non-local interactions may contribute to stabilizing the secondary structure. We thus investigated whether such non-local interactions included intramolecular contacts between residues in Tau-1 and Tau-5, similarly to what has been shown to occur in globular proteins under mild

denaturing conditions³⁷. For this analysis we used an NTD construct encompassing the part of Tau-5 contained in AF-1*, corresponding to residues 330-448 (termed Tau-5*, Fig. 1b). We compared the ^1H and ^{15}N chemical shifts of Tau-5* to those of AF-1* and observed that they were highly similar (Fig. S2). These results indicate that the chemical environment that residues in the Tau-5 region experience is the same both in the presence and absence of Tau-1. We conclude that the secondary structure present in AF-1* does not rely on long-range interactions between residues in Tau-1 and residues in Tau-5.

EPI-001 interacts reversibly with Tau-5 It has been proposed that the mechanism of action of EPI-001 (Fig. 3a) involves two steps¹³. The first step being the formation of a reversible complex between this compound and a specific conformation of AF-1 and the second the nucleophilic attack, by a protein side chain, on the C-Cl bond of EPI-001 to form an adduct unable to activate transcription. To investigate the first step of this mechanism and characterize the putative reversible complex we synthesized EPI-001 (details available as SI) and used NMR to analyze its effect on the resonances of AF-1* at 278 K. As shown by MS under these conditions the irreversible reaction of EPI-001 with AF-1* is sufficiently slow to allow the study of the reversible interaction by NMR (Fig. S5).

The results that we obtained, shown in figures 3b and c, indicate that EPI-001 causes small but highly reproducible changes in ^{15}N chemical shifts in residues 354 to 448 of AF-1*, that correspond to Tau-5. Interaction with EPI-001 affects the resonances of a large number of residues, which are found in the three regions of sequence in Tau-5 identified as partially folded by the combined analysis of the backbone ^{13}C chemical shifts and ^{15}N transverse relaxation rates (Figs. 2b, 2d and 3b). We detected much smaller perturbations of the resonances of residues corresponding to Tau-1 even though this region contains partially helical regions such as residues in core Tau-1 (Figs. 2b and 3b). These results indicate that EPI-001 does not simply interact with any region of sequence with helical propensity but, rather, that this compound targets Tau-5 due to the ability of this sub-domain to adopt a conformation or, most likely, an ensemble of conformations that have affinity for it.

1
2
3
4
5 With the aim of further investigating the nature of the interaction we monitored the effect
6 of AF-1* on the resonances of EPI-001. We observed that sub-stoichiometric amounts
7 (25 μ M and 50 μ M) of AF-1* caused small but reproducible chemical shift changes in the
8 1 H NMR spectrum of EPI-001 at 250 μ M, as well as increases in linewidth similar to
9 those commonly observed in small drug-like molecules transiently associating with
10 macromolecules (Fig. S3). Together with our observation of perturbations in a large
11 number of residues of AF-1* this result suggests that EPI-001 interacts with an
12 ensemble of conformations adopted by AF-1* where these regions of sequence adopt a
13 partially folded structure.
14
15
16
17
18
19

20
21 A large number of residues in Tau-5 experience chemical shift perturbations, more than
22 what is expected for the binding of a small molecule. This can be due to the inspecific
23 interaction of one or various molecules of EPI-001 with three independent interaction
24 sites or to the combination of direct and indirect chemical shift changes due to structural
25 changes occurring in Tau-5 upon interaction with EPI-001³⁸. To exclude that one or more
26 EPI-001 molecules interact independently with the three regions of sequence of AF-1*
27 where chemical shift perturbations are observed we monitored the effect on its 1 H NMR
28 spectrum of adding three peptides (R1 to R3) with sequences corresponding to regions
29 341-371, 391-414 and 426-446, respectively (details available as SI). We found that
30 these peptides caused no changes in the spectrum of EPI-001, indicating that EPI-001
31 does not interact with them (Fig. S3) and confirming that residues 354-448 must
32 simultaneously be present because they are necessary for binding this compound or for
33 stabilizing the bound state of Tau-5* (Fig. 4).
34
35
36
37
38
39
40
41
42

43
44 It is interesting to note that Tau-5, the region of sequence of the NTD targeted by EPI-
45 001, is partially folded, but it is important to emphasize that the data that we have
46 obtained do not provide us with a mechanism of molecular recognition. It is possible that
47 a conformational selection mechanism operates *i.e.* that EPI-001 interacts with a subset
48 of the conformations that Tau-5* samples in equilibrium, but also that EPI-001 induces a
49 new conformation in Tau-5*. Further work will be necessary to differentiate these two
50 scenarios.
51
52
53
54
55
56
57
58
59
60

EPI-001 has two stereogenic centers and can therefore be found as four stereoisomers. To investigate whether the interaction between this compound and Tau-5 is stereospecific we synthesized the four stereoisomers (details available as SI) and studied their interaction with AF-1* by NMR. The results that we obtained (Fig. S4) indicate that the four compounds can interact with the NTD of AR, and thus the interaction appears to occur with little or no stereoselectivity. These results are in agreement with results obtained *in vivo* by Myung *et al*, who found that, although one of the stereoisomers tested was slightly more active than the other ones, the inhibitory activity of the four stereoisomers was similar.¹³

Put together our results indicate that AF-1 is partially folded in regions of sequence that correspond to those which are functionally relevant for interacting with the transcription machinery and co-regulators of transcription. In addition, they reveal that Tau-1 and Tau-5, the two independent transcription activation units that are found in AF-1, correspond to different sub-domains that appear not to be involved in long-range interactions *i.e.* are structurally and dynamically, at least under our conditions, independent. Finally, and most importantly, they show that Tau-5, that plays a particularly important role in AR activation in the absence of androgens, can be targeted by compounds such as EPI-001 (Fig. 4). Although the lack of stereoselectivity that we observe suggests that the binding mode of EPI-001 may not be sufficiently well-defined for conventional drug development our results are of relevance for drug discovery for CRPC because they suggests that the NTD, and Tau-5 in particular, may represent a suitable therapeutic target.

ACKNOWLEDGEMENTS

We thank J. M. Valverde (IRB) as well as the NMR facilities of the University of Barcelona (CCiT UB) and the Instituto de Química Física Rocasolano (IQFR, CSIC) for their assistance in, respectively, protein production and NMR. This work was supported by IRB, ICREA (X.S.), Obra Social “la Caixa” (Fellowship to E.D.M. and CancerTec grants to X.S.) MICINN (CTQ2009-08850 to X.S.), MINECO (BIO2012-31043 to X.S.; CTQ2014-56361-P to A.R), Marató de TV3 (102030 to X.S. and 102031 to E.E.P) the COFUND programme of the European Commission (C.T.W.P., A. R. and X.S.), the European Research Council (CONCERT, contract number 648201, to X.S.), the Ramón y Cajal program of MICINN (RYC-2011-07873 to C.W.B.) the Serra Hunter Programme

(E.E.P.) and AGAUR (SGR-2014-56RR14 to E.E.P). IRB Barcelona is the recipient of a Severo Ochoa Award of Excellence from MINECO (Government of Spain).

Supporting Information Available: This material is available free of charge *via* the Internet

FIGURE LEGENDS

Figure 1 Predicted properties of the sequence of the transactivation domain of AR a) Domain organization of AR^{34,39} with an indication of the position of Zn atoms in the DBD (grey) and of dihydrotestosterone (DHT) in the LBD (blue). b) Definitions of activation function 1 (AF-1), transcription activation units 1 and 5 (Tau-1 and Tau-5), AF-1* and Tau-5* (the regions of sequence studied in this work) with an indication of the regions of low sequence complexity such as polyGln (Q_n), polyPro (P_n), polyAla (A_n) and polyGly (G_n) tracts. c) Propensity to disorder of the NTD predicted by PONDR VL-XT²⁶ with an indication of the functional motifs defining the core of Tau-1 and Tau-5, shaded in red and of the region of sequence studied in this work, shaded in grey. d) Positions of the motifs of the NTD of AR involved in protein protein interactions and acronyms of the binding partners (see main text for details). e) Propensity to adopt α -helical conformations predicted by Agadir³⁶, as a function of residue number, with an indication of the core of Tau-1 and Tau-5 (shaded in red) and AF-1* (shaded in grey).

Figure 2 Structural properties of the transactivation domain revealed by solution NMR a) ¹H, ¹⁵N-HSQC spectrum of AF-1*. b) Plot of the difference between the secondary C α and C β chemical shifts measured for AF-1* with an indication of the regions of sequence with predicted disorder propensity (DP) lower than 50%, in grey, with predicted helical propensity (hel) higher than 10%, in blue, and involved in protein protein interactions (PPIs), in green and, at the bottom of the top panel, an indication of the nascent secondary structure identified experimentally (SS, cylinders indicate helical propensity, black rectangles indicate propensity to adopt an extended conformation). c) Plot of the secondary structure propensity of the residues of AF-1*, where SSP=1 indicates a fully formed helix and SSP=1 an extended conformation, obtained by using the algorithm SSP to extract the information on secondary structure contained in backbone (¹³C α , ¹³C β , ¹³CO, ¹⁵N, HN) chemical shifts. d) Plot of the ¹⁵N transverse relaxation rates (R₂) of the residues of AF-1* at 250 μ M. The cores of Tau-1 and Tau-5 are shaded in red.

Figure 3 EPI-001 selectively interacts with transcription activation unit 5 of the transactivation domain of AR a) Structure of EPI-001 with an indication of the two stereogenic centres with the symbol *. b) Plot, as a function of residue number, of the

1
2
3 change in ^{15}N chemical shift of AF-1* caused by addition of EPI-001. c) Selected regions
4 of the ^1H , ^{15}N -HSQC spectrum of 25 μM AF-1* in the absence (blue) and in the presence
5 (red) of 10 molar equivalents of EPI-001.
6
7
8

9
10 **Figure 4** Scheme of the interaction of EPI-001 with partially folded of Tau-5.
11
12
13
14
15
16
17
18
19
20
21
22
23
24
25
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60

REFERENCES

- (1) Gelmann, E. P. (2002) Molecular Biology of the Androgen Receptor. *J. Clin. Oncol.* 20, 3001–3015.
- (2) Akaza, H., Hinotsu, S., Usami, M., Arai, Y., Kanetake, H., Naito, S., and Hirao, Y. (2009) Combined androgen blockade with bicalutamide for advanced prostate cancer. *Cancer* 115, 3437–3445.
- (3) Harris, W. P., Mostaghel, E. A., Nelson, P. S., and Montgomery, B. (2009) Androgen deprivation therapy: progress in understanding mechanisms of resistance and optimizing androgen depletion. *Nat. Clin. Pract. Urol.* 6, 76–85.
- (4) Karantanos, T., Corn, P. G., and Thompson, T. C. (2013) Prostate cancer progression after androgen deprivation therapy: mechanisms of castrate resistance and novel therapeutic approaches. *Oncogene* 32, 5501–5511.
- (5) Reid, J., Kelly, S. M., Watt, K., Price, N. C., and McEwan, I. J. (2002) Conformational analysis of the androgen receptor amino-terminal domain involved in transactivation. Influence of structure-stabilizing solutes and protein-protein interactions. *J. Biol. Chem.* 277, 20079–20086.
- (6) Lavery, D. N., and McEwan, I. J. (2008) Structural characterization of the native NH₂-terminal transactivation domain of the human androgen receptor: a collapsed disordered conformation underlies structural plasticity and protein-induced folding. *Biochemistry* 47, 3360–3369.
- (7) Quayle, S. N., Mawji, N. R., Wang, J., and Sadar, M. D. (2007) Androgen receptor decoy molecules block the growth of prostate cancer. *Proc. Natl. Acad. Sci. U. S. A.* 104, 1331–1336.
- (8) Lamberto, G. R., Torres-Monserrat, V., Bertoncini, C. W., Salvatella, X., Zweckstetter, M., Griesinger, C., and Fernandez, C. O. (2011) Toward the discovery of effective polycyclic inhibitors of alpha-synuclein amyloid assembly. *J. Biol. Chem.* 286, 32036–32044.
- (9) Krishnan, N., Koveal, D., Miller, D. H., Xue, B., Akshinthala, S. D., Kragelj, J., Jensen, M. R., Gauss, C.-M., Page, R., Blackledge, M., Muthuswamy, S. K., Peti, W., and Tonks, N. K. (2014) Targeting the disordered C terminus of PTP1B with an allosteric inhibitor. *Nat. Chem. Biol.* 10, 558–566.
- (10) Metallo, S. J. (2010) Intrinsically disordered proteins are potential drug targets. *Curr. Opin. Chem. Biol.* 14, 481–488.
- (11) Heller, G. T., Sormanni, P., and Vendruscolo, M. (2015) Targeting disordered proteins with small molecules using entropy. *Trends Biochem. Sci.* 40, 491–496.
- (12) Andersen, R. J., Mawji, N. R., Wang, J., Wang, G., Haile, S., Myung, J.-K., Watt, K., Tam, T., Yang, Y. C., Bañuelos, C. A., Williams, D. E., McEwan, I. J., Wang, Y., and Sadar, M. D. (2010) Regression of castrate-recurrent prostate cancer by a small-molecule inhibitor of the amino-terminus domain of the androgen receptor. *Cancer Cell* 17, 535–546.
- (13) Myung, J.-K., Bañuelos, C. A., Fernandez, J. G., Mawji, N. R., Wang, J., Tien, A. H., Yang, Y. C., Tavakoli, I., Haile, S., Watt, K., McEwan, I. J., Plymate, S., Andersen, R. J., and Sadar, M. D. (2013) An androgen receptor N-terminal domain antagonist for treating prostate cancer. *J. Clin. Invest.* 123, 2948–2960.
- (14) Neidhardt, F. C., Bloch, P. L., and Smith, D. F. (1974) Culture medium for enterobacteria. *J. Bacteriol.* 119, 736–747.
- (15) McEwan, I. J. (2012) Intrinsic disorder in the androgen receptor: identification, characterisation and drugability. *Mol. Biosyst.* 8, 82–90.

- (16) Dunker, A. K., Lawson, J. D., Brown, C. J., Williams, R. M., Romero, P., Oh, J. S., Oldfield, C. J., Campen, A. M., Ratliff, C. M., Hipps, K. W., Ausio, J., Nissen, M. S., Reeves, R., Kang, C., Kissinger, C. R., Bailey, R. W., Griswold, M. D., Chiu, W., Garner, E. C., and Obradovic, Z. (2001) Intrinsically disordered protein. *J. Mol. Graph. Model.* **19**, 26–59.
- (17) Mohan, A., Oldfield, C. J., Radivojac, P., Vacic, V., Cortese, M. S., Dunker, A. K., and Uversky, V. N. (2006) Analysis of molecular recognition features (MoRFs). *J. Mol. Biol.* **362**, 1043–1059.
- (18) Jenster, G., van der Korput, H. A., Trapman, J., and Brinkmann, A. O. (1995) Identification of two transcription activation units in the N-terminal domain of the human androgen receptor. *J. Biol. Chem.* **270**, 7341–7346.
- (19) Callewaert, L., Van Tilborgh, N., and Claessens, F. (2006) Interplay between two hormone-independent activation domains in the androgen receptor. *Cancer Res.* **66**, 543–553.
- (20) Dehm, S. M., Regan, K. M., Schmidt, L. J., and Tindall, D. J. (2007) Selective role of an NH₂-terminal WxxLF motif for aberrant androgen receptor activation in androgen depletion independent prostate cancer cells. *Cancer Res.* **67**, 10067–10077.
- (21) Metzger, E., Müller, J. M., Ferrari, S., Buettner, R., and Schüle, R. (2003) A novel inducible transactivation domain in the androgen receptor: implications for PRK in prostate cancer. *EMBO J.* **22**, 270–280.
- (22) Kumar, R., Betney, R., Li, J., Thompson, E. B., and McEwan, I. J. (2004) Induced alpha-helix structure in AF1 of the androgen receptor upon binding transcription factor TFIIF. *Biochemistry* **43**, 3008–3013.
- (23) Tamiola, K., Acar, B., and Mulder, F. A. A. (2010) Sequence-specific random coil chemical shifts of intrinsically disordered proteins. *J. Am. Chem. Soc.* **132**, 18000–18003.
- (24) Marsh, J. A., Singh, V. K., Jia, Z., and Forman-Kay, J. D. (2006) Sensitivity of secondary structure propensities to sequence differences between alpha- and gamma-synuclein: implications for fibrillation. *Protein Sci.* **15**, 2795–2804.
- (25) Klein-Seetharaman, J., Oikawa, M., Grimshaw, S. B., Wirmer, J., Duchardt, E., Ueda, T., Imoto, T., Smith, L. J., Dobson, C. M., and Schwalbe, H. (2002) Long-range interactions within a nonnative protein. *Science* **295**, 1719–1722.
- (26) Romero, P., Obradovic, Z., Li, X., Garner, E. C., Brown, C. J., and Dunker, A. K. (2001) Sequence complexity of disordered protein. *Proteins: Struct. Funct. Bioinf.* **42**, 38–48.
- (27) Zhu, P., Baek, S. H., Bourk, E. M., Ohgi, K. A., Garcia-Bassets, I., Sanjo, H., Akira, S., Kotol, P. F., Glass, C. K., Rosenfeld, M. G., and Rose, D. W. (2006) Macrophage/cancer cell interactions mediate hormone resistance by a nuclear receptor derepression pathway. *Cell* **124**, 615–629.
- (28) Asangani, I. A., Dommeti, V. L., Wang, X., Malik, R., Cieslik, M., Yang, R., Escara-Wilke, J., Wilder-Romans, K., Dhanireddy, S., Engelke, C., Iyer, M. K., Jing, X., Wu, Y.-M., Cao, X., Qin, Z. S., Wang, S., Feng, F. Y., and Chinnaiyan, A. M. (2014) Therapeutic targeting of BET bromodomain proteins in castration-resistant prostate cancer. *Nature* **510**, 278–282.
- (29) He, B., Bai, S., Hnat, A. T., Kalman, R. I., Minges, J. T., Patterson, C., and Wilson, E. M. (2004) An androgen receptor NH₂-terminal conserved motif interacts with the COOH terminus of the Hsp70-interacting protein (CHIP). *J. Biol. Chem.* **279**, 30643–30653.
- (30) Migliaccio, A., Castoria, G., Giovannelli, P., and Auricchio, F. (2010) Cross talk between epidermal growth factor (EGF) receptor and extra nuclear steroid receptors in cell lines. *Mol. Cell. Endocrinol.* **327**, 19–24.

- (31) Gioeli, D., Ficarro, S. B., Kwiek, J. J., Aaronson, D., Hancock, M., Catling, A. D., White, F. M., Christian, R. E., Settlage, R. E., Shabanowitz, J., Hunt, D. F., and Weber, M. J. (2002) Androgen receptor phosphorylation. Regulation and identification of the phosphorylation sites. *J. Biol. Chem.* 277, 29304–29314.
- (32) Adzhubei, A. A., Sternberg, M. J. E., and Makarov, A. A. (2013) Polyproline-II helix in proteins: structure and function. *J. Mol. Biol.* 425, 2100–2132.
- (33) He, B., Kemppainen, J. A., and Wilson, E. M. (2000) FXXLF and WXXLF sequences mediate the NH2-terminal interaction with the ligand binding domain of the androgen receptor. *J. Biol. Chem.* 275, 22986–22994.
- (34) Hur, E., Pfaff, S. J., Payne, E. S., Grøn, H., Buehrer, B. M., and Fletterick, R. J. (2004) Recognition and accommodation at the androgen receptor coactivator binding interface. *PLoS Biol.* 2, E274.
- (35) Lagarde, W. H., Blackwelder, A. J., Mingos, J. T., Hnat, A. T., French, F. S., and Wilson, E. M. (2012) Androgen receptor exon 1 mutation causes androgen insensitivity by creating phosphorylation site and inhibiting melanoma antigen-A11 activation of NH2- and carboxyl-terminal interaction-dependent transactivation. *J. Biol. Chem.* 287, 10905–10915.
- (36) Muñoz, V., and Serrano, L. (1994) Elucidating the folding problem of helical peptides using empirical parameters. *Nat. Struct. Biol.* 1, 399–409.
- (37) Bruun, S. W., Iešmantavičius, V., Danielsson, J., and Poulsen, F. M. (2010) Cooperative formation of native-like tertiary contacts in the ensemble of unfolded states of a four-helix protein. *Proc. Natl. Acad. Sci. U. S. A.* 107, 13306–13311.
- (38) Fenwick, R. B., Esteban-Martín, S., and Salvatella, X. (2011) Understanding biomolecular motion, recognition, and allostery by use of conformational ensembles. *Eur. Biophys. J.* 40, 1339–1355.
- (39) Shaffer, P. L., Jivan, A., Dollins, D. E., Claessens, F., and Gewirth, D. T. (2004) Structural basis of androgen receptor binding to selective androgen response elements. *Proc. Natl. Acad. Sci. U. S. A.* 101, 4758–4763.

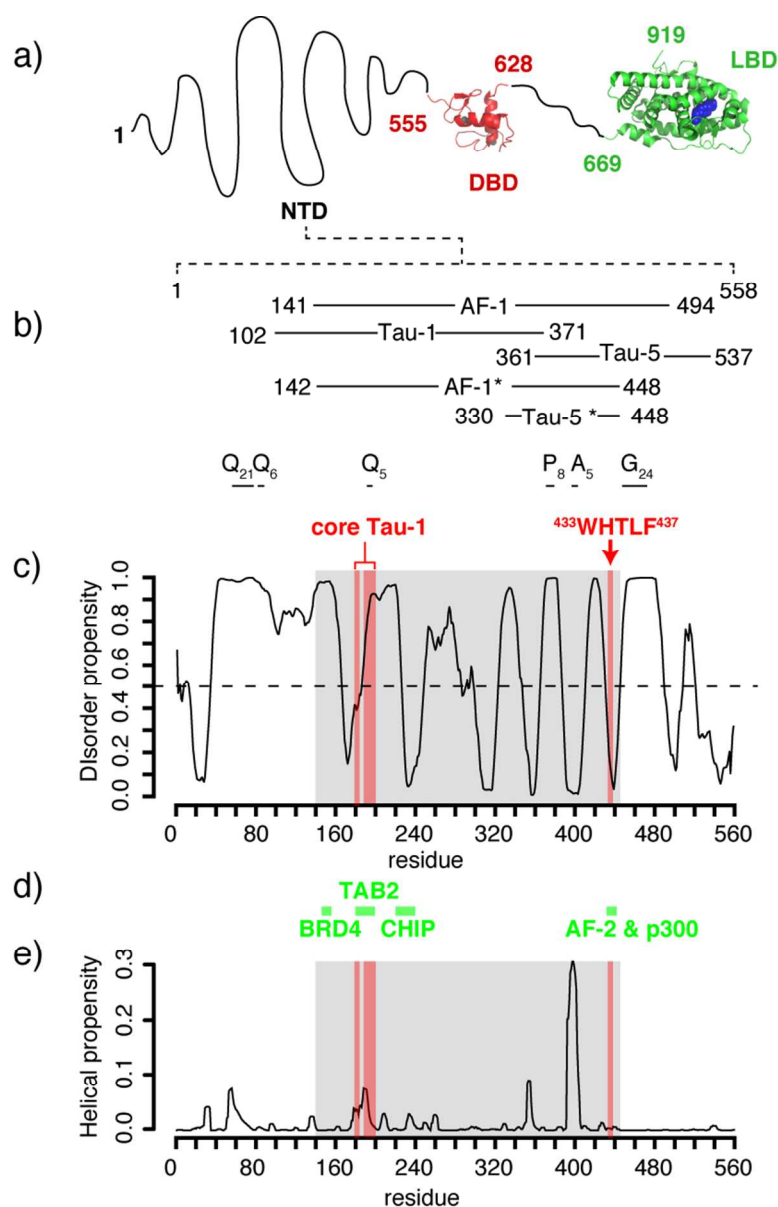


Figure 1 Predicted properties of the sequence of the transactivation domain of AR
70x109mm (300 x 300 DPI)

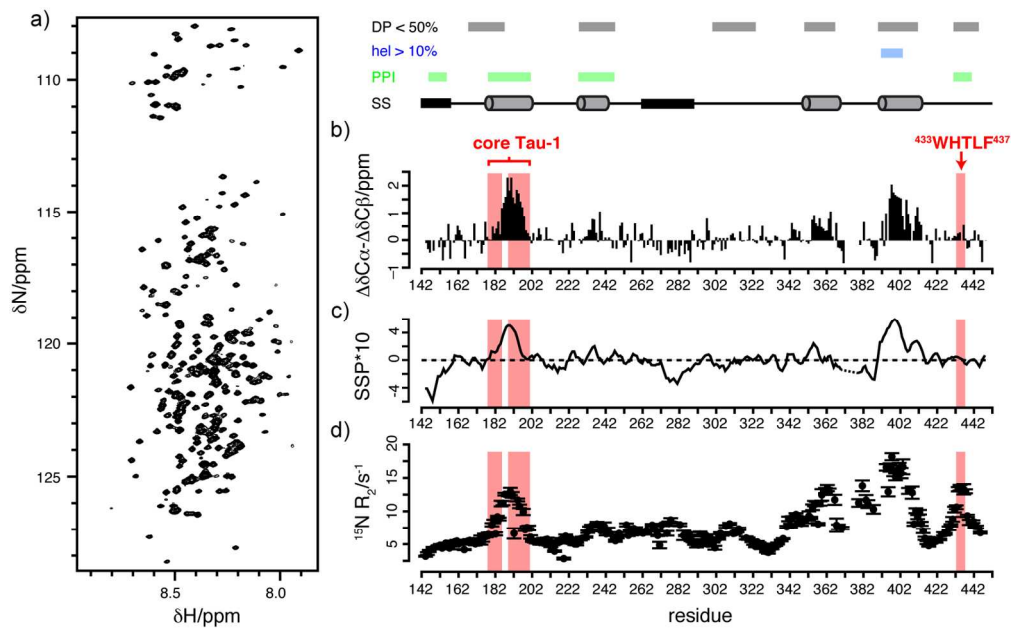


Figure 2 Structural properties of the transactivation domain revealed by solution NMR
144x90mm (300 x 300 DPI)

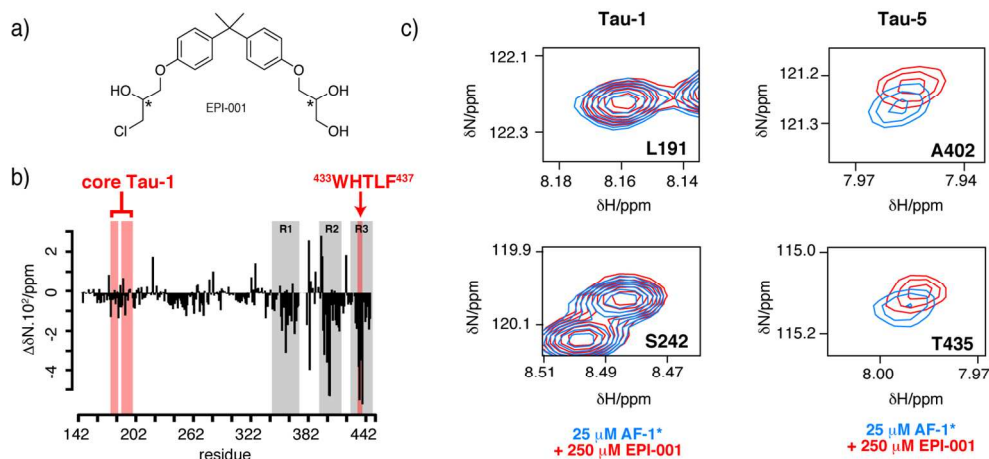


Figure 3 EPI-001 selectively interacts with transcription activation unit 5 of the transactivation domain of AR 133x61mm (300 x 300 DPI)

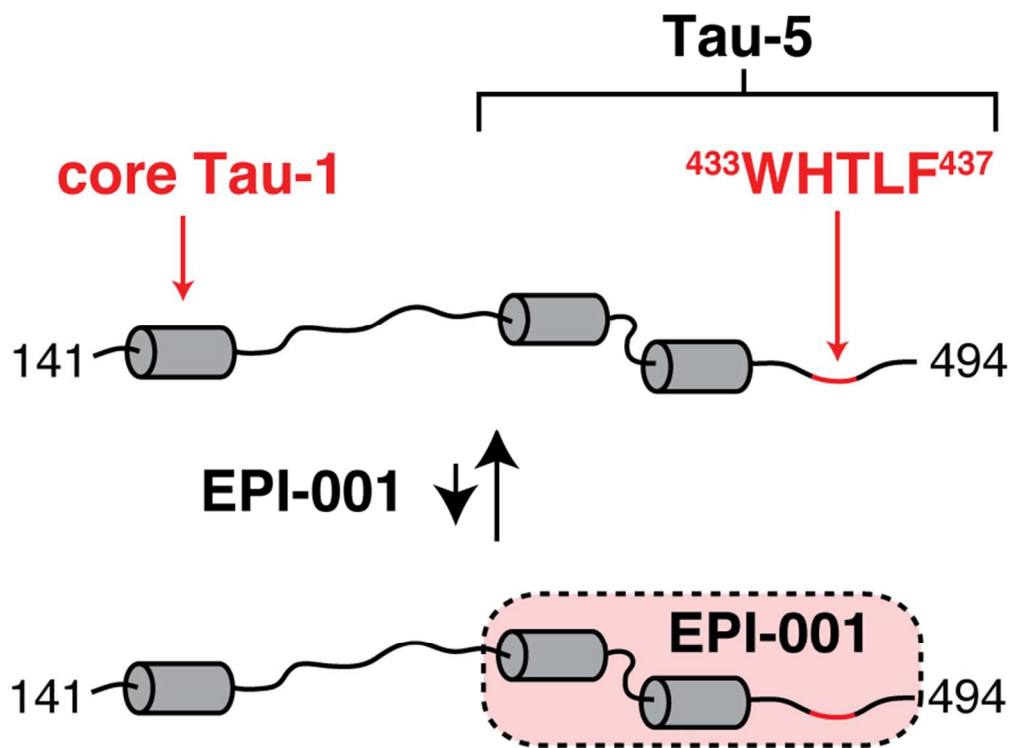
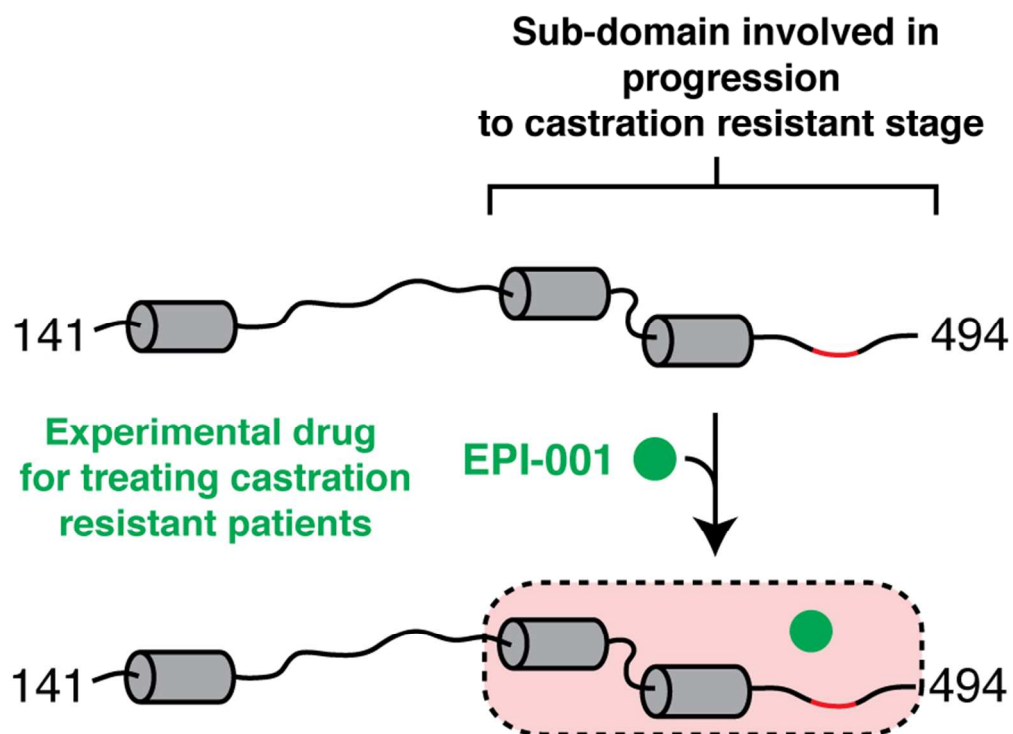


Figure 4 Scheme of the interaction of EPI-001 with partially folded of Tau-5
74x55mm (300 x 300 DPI)



TOC figure
74x53mm (300 x 300 DPI)